



# **STIC Search Report**

## **Biotech-Chem Library**

**STIC Database Tracking Number: 139767**

**TO: James Schultz**  
**Location: REM/2D18/2C18**  
**Art Unit: 1635**  
**Wednesday, December 08, 2004**

**Case Serial Number: 09/913800**

**From: David Schreiber**  
**Location: Biotech-Chem Library**  
**Remsen E01A61**  
**Phone: 272-2526**

**david.schreiber@uspto.gov**

### **Search Notes**

# SEARCH REQUEST FORM

Scientific and Technical Information Center

Requester's Full Name: \_\_\_\_\_ Examiner #: \_\_\_\_\_ Date: \_\_\_\_\_  
 Art Unit: \_\_\_\_\_ Phone Number 30 \_\_\_\_\_ Serial Number: \_\_\_\_\_  
 Mail Box and Bldg/Room Location: \_\_\_\_\_ Results Format Preferred (circle): PAPER DISK E-MAIL

**If more than one search is submitted, please prioritize searches in order of need.**

\*\*\*\*\*

Please provide a detailed statement of the search topic, and describe as specifically as possible the subject matter to be searched. include the elected species or structures, keywords, synonyms, acronyms, and registry numbers, and combine with the concept or utility of the invention. Define any terms that may have a special meaning. Give examples or relevant citations, authors, etc, if known. Please attach a copy of the cover sheet, pertinent claims, and abstract.

Title of invention: \_\_\_\_\_

Inventors (please provide full names): \_\_\_\_\_

Earliest Priority Filing Date: \_\_\_\_\_

*\*For Sequence Searches Only\* Please include all pertinent information (parent, child, divisional, or issued patent numbers) along with the appropriate serial number.*

\*\*\*\*\*

**STAFF USE ONLY**

	Type of Search	Vendors and cost where applicable
Searcher: <u>P. Schreiber</u>	NA Sequence (#) <u>22</u>	STN _____
Searcher Phone #: <u>272-2526</u>	AA Sequence (#) _____	Dialog _____
Searcher Location: <u>Ramsen E01A61</u>	Structure (#) _____	Questel/Orbit _____
Date Searcher Picked Up _____	Bibliographic _____	Dr.Link _____
Date Completed: <u>12/8</u>	Litigation _____	Lexis/Nexis _____
Searcher Prep. Review Time <u>15</u>	Fulltext _____	Sequence Systems <u>Compuer</u>
Clerical Prep. Time: _____	Patent Family _____	WWW/Internet _____
Online Time <u>126</u>	Other _____	Other (specify) _____

Schreiber, David

139767

**From:** Schultz, James  
**Sent:** Wednesday, November 17, 2004 10:09 AM  
**To:** Schreiber, David  
**Subject:** Score over length search 09/913,800

Hi David,

I need a score over length nucleotide sequence search on SEQ ID NOS: 32 and 21 in the above entitled case. I need the lower and upper limits to be 8 and 30, respectively, I only need hits that are 100% complementary, and please transfer as many hits into the excel program as possible. No need to search the interference databases at this time.

I realize you don't normally search more than two sequences, but applicants have asserted that these two sequences are usable together, and that they both comprise the invention...I don't see any way around searching these.

Also, can you give me an idea how long it is typically taking to get results back these days for this type of search?

Thanks,  
Doug Schultz

*James Douglas Schultz, PhD*  
AU 1635 (Biotechnology)  
Patent Examiner  
United States Patent and Trademark Office  
(Office) REM 2D18  
(Mail) REM 2C18  
(571) 272-0763

## SCORE OVER LENGTH SEARCHES

Attached is a score over length search. This search was developed to overcome limitations in most standard search systems which favor large sequences with high scoring, but lesser overall identity over smaller sequences with higher overall identity. This search is especially useful for relatively small nucleic acid or polypeptide target sequences (antisense, fragments, probes, primers, RNAi, epitopes, haptens, etc.) claimed functionally via a form of hybridization and/or identity language and having defined upper and lower polynucleotide and or polypeptide length limits.

The score over length search is performed by first running the query sequence using examiner-specified identity and polynucleotide or protein length limit parameters, and saving 65,000 hits and 0 alignments from each desired database. The resulting output is reformatted using a Microsoft Word macro and is imported into Excel. The summary table data are then sorted by the ratio of score of each hit sequence divided by its length and the accession numbers for all hits below the examiner's desired score over length parameters are deleted. The remaining accession numbers are used to pull the corresponding sequences from the databases into subdatabases enriched for good hits and the query sequence is re-run against these subdatabases to yield the final results.

The score over length cutoff for this search is 100%.

Examiner Please Note: This cover sheet should be included when submitting results to be scanned.

GenCore version 5.1.6  
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OM nucleic - nucleic search, using SW model

Run on: December 8, 2004, 06:35:33 ; Search time 0.001 Seconds  
(without alignments)  
1,908 Million cell updates/sec

Title: US-09-913-800-21  
Perfect score: 18  
Sequence: 1 cctgagcctgttgcgcac 18

Scoring table: IDENTITY NUC  
Gapop 10.0, Gapext 0.5

Searched: 5 seqs, 53 residues  
Total number of hits satisfying chosen parameters: 10

Minimum DB seq length: 8  
Maximum DB seq length: 30

Post-Processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 5 summaries

Database: rng21.seq.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	18	100.0	18	AAA08468	Human Akt-2 phosph
2	9	50.0	9	AB072180	Zinc finger protei
3	9	50.0	9	ADA64507	Zinc finger target
4	9	50.0	9	ADM23199	Synthetic zinc fin
5	8	44.4	8	AAA80787	A. thaliana primer

ALIGNMENTS

RESULT 1  
AAA08468 standard; DNA; 18 BP.  
AAA08468;  
17-JUL-2000 (first entry)  
Human Akt-2 phosphorothioate antisense oligonucleotide SEQ ID NO:21.  
Human, Akt-2; antisense oligonucleotide; phosphorothioate; inhibition;  
serine/threonine kinase; antiinflammatory; cytoskeletal; antileukemic;  
gene therapy; infection; inflammation; tumour; ss.  
Homo sapiens.  
Location/Qualifiers  
Key modified\_base 1..18 a  
/\*tag= a  
/note= "phosphorothioate linkages"  
US6043090-A.

28-MAR-2000.  
23-FEB-1999; 99US-00256465.  
23-FEB-1999; 99US-00256465.  
(ISIS-) ISIS PHARM INC.  
Monla BP, Cowsett LM;  
WPI; 2000-270345/23.  
Antisense compound for diagnosis and treatment of infection, inflammation and tumor formation is targeted towards the nucleic acid encoding a member of serine/threonine family of kinases.

Claim 3; Col 38; 30pd; English.  
The present invention describes antisense compounds of about 8-30 nucleotides in length targeted to the 5' UTR (untranslated region), 3' UTR or coding region of the nucleic acid encoding human Akt-2, which inhibits the expression of human Akt-2. Human Akt-2 is a member of the Akt/PKB family of serine/threonine kinases. The antisense compounds have antiinflammatory, cyostatic and antileukemic activities, and can be used in gene therapy. They are useful in inhibiting the expression of human Akt-2 by contacting the cells or the tissues in vitro. They can also be used for diagnosis and treatment of infection, inflammation and tumor formation, and for prophylaxis. The present sequence represents a human Akt-2 phosphorothioate antisense oligonucleotide used in the exemplification of the present invention

Sequence 18 BP; 2 A; 4 C; 7 G; 5 T; 0 U; 0 Other;  
Query Match 100.0%; Score 18; DB 1; Length 18;  
Best Local Similarity 100.0%; Pred. No. 0;  
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 CTTGAGGCTGTGGCGAC 18  
DB 1 CTTGAGGCTGTGGCGAC 18

RESULT 2  
AB072180  
ID ABQ72180 standard; DNA; 9 BP.  
ABQ72180;  
28-AUG-2002 (first entry)  
Zinc finger protein related oligonucleotide target SEQ ID NO:2478.  
Zinc finger protein; ZFP; DNA binding protein; zinc finger; ss.  
Homo sapiens.  
Synthetic.  
WO200242459-A2.  
30-MAY-2002.  
20-NOV-2001; 2001WO-US043438.  
20-NOV-2000; 2000US-00716637.  
(SANG-) SANGAMO BIOSCIENCES INC.  
Liu Q;  
WPI; 2002-500284/53.  
New zinc finger protein that binds to target site, useful in studying gene function and for human therapeutics and plant engineering, comprises

PT first, second and third zinc fingers, ordered from N- to C-terminus.  
PS Example 1; Page 63; 81pp; English.

CC The present invention describes a zinc finger protein (I) that binds to a  
CC target site, comprising a first (F1), a second (F2), and a third (F3)  
CC zinc finger, ordered F1, F2, F3 from N-terminus to C-terminus, where the  
CC target site comprises, in 3'-5' direction, a first (S1), a second (S2),  
CC and a third (S3) target sub-site. Also described are: (1) a polypeptide  
CC (II) comprising (I); (2) a polynucleotide (III) encoding (I) or (II); and  
CC (3) designing (M) (I) involves selecting the F1 zinc finger such that it  
CC binds to the S1 target sub-site, selecting the F2 zinc finger such that it  
CC binds to the S2 target sub-site, and selecting the F3 zinc finger such  
CC that it binds to the S3 target sub-site, thus designing (I) that binds to  
CC a target site. (I) is useful for recognition of triplet target sub-sites  
CC having the nucleotide G in the 5'-most position of the sub-site. (I) is  
CC useful in studying gene function, and for human therapeutics and plant  
CC engineering. (II), (III) or (III) is useful in therapeutic methods to  
CC modulate the expression of a target region within a subject, in  
CC diagnostic methods for sequence specific detection of target nucleic acid  
CC in a sample, and in assays to determine the phenotype and function of  
CC gene expression. (II) has improved affinity and specificity for their  
CC target sequences, as well as enhanced biological activity. ABQ71213 to  
CC ABQ72214 and ABP48191 to ABP51230 represent DNA target sequences and zinc  
CC finger peptides which are given in the exemplification of the present  
CC invention

CC Sequence 9 BP; 1 A; 1 C; 4 G; 3 T; 0 U; 0 Other;

Query Match 50.0%; Score 9; DB 1; Length 9;  
Best Local Similarity 100.0%; Pred. No. 0;  
Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 4 GAGGCTGTT 12  
|||  
1 GAGGCTGTT 9

RESULT 3  
ADA64507

ID ADA64507 standard; DNA; 9 BP.

AC ADA64507;

XX 20-NOV-2003 (first entry)

DE Zinc finger target sequence DNA #965.

KW ds: target sequence; zinc finger protein; improved affinity;  
KW multi-finger zinc finger protein; improved affinity;  
KW improved specificity; enhanced biological activity.

OS Synthetic.

PN US2003068675-A1.

PD 10-APR-2003.

PF 20-NOV-2001; 2001US-00990186.

PR 24-MAR-1999; 99US-0126238P.

PR 24-MAR-1999; 99US-0126239P.

PR 30-JUL-1999; 99US-0146595P.

PR 30-JUL-1999; 99US-0146615P.

PR 23-MAR-2000; 2000US-00535008.

PR 20-NOV-2000; 2000US-00716637.

PA (LIUQ/) LIU Q.

PI Liu Q;

XX WPI; 2003-567233/53.

PT Designing zinc finger protein that has three zinc fingers from N-terminus  
PT and C-terminus that bind to sub-sites in 3' to 5' direction, in a target  
PT site, by selecting zinc fingers that bind their respective sub-sites.  
XX

PS Disclosure; Page 27; 34pp; English.

CC The invention relates to a method of designing a zinc finger protein. The  
CC method is useful for designing a zinc finger protein. The method provides  
CC multi-finger zinc finger proteins with improved affinity and specificity  
CC for their target sequences, as well as enhanced biological activity. The  
CC present sequence represents a zinc finger protein DNA target sequence.

CC Sequence 9 BP; 1 A; 1 C; 4 G; 3 T; 0 U; 0 Other;

Query Match 50.0%; Score 9; DB 1; Length 9;  
Best Local Similarity 100.0%; Pred. No. 0;  
Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 4 GAGGCTGTT 12  
|||  
1 GAGGCTGTT 9

RESULT 4

ID ADM23199 standard; DNA; 9 BP.

AC ADM23199;

XX 20-MAY-2004 (first entry)

DE Synthetic zinc finger protein target DNA #965.

KW zinc finger protein; triplet target sub-site; zinc finger motif; sp-1; ds.  
KW Unidentified.

OS Unidentified.

PN US2003104526-A1.

PD 05-JUN-2003.

PF 20-NOV-2001; 2001US-00989994.

PR 24-MAR-1999; 99US-0126238P.

PR 24-MAR-1999; 99US-0126239P.

PR 30-JUL-1999; 99US-0146595P.

PR 30-JUL-1999; 99US-0146615P.

PR 23-MAR-2000; 2000US-00535008.

PR 20-NOV-2000; 2000US-00716637.

PA (LIUQ/) LIU Q.

PI Liu Q;

XX WPI; 2003-843091/78.

PT New zinc finger protein used for recognizing triplet target sub-sites  
PT having nucleotide G in 5'-most position of sub-site, that has been  
PT optimized with respect to location of sub-site within target site.  
XX

PS Example 6; SEQ ID NO 2478; 48pp; English.

CC The invention describes a new zinc finger protein that binds to a target  
CC site comprising a first (F1), a second (F2) or a third (F3) zinc finger,  
CC ordered F1, F2 and F3 from N-terminus to C-terminus. The target site  
CC comprises, in the 3' to 5' direction, first (S1), second (S2) and third  
CC (S3) target sub-sites. The zinc finger proteins can be used for  
CC recognizing triplet target sub-sites having the nucleotide G in the 5'-  
CC most position of the sub-site, that has been optimised with respect to the  
CC location of the sub-site within the target site. This sequence represents  
CC the target polynucleotide to which the zinc finger protein sp-1 consensus  
CC sequence binds.

SQ Sequence 9 BP; 1 A; 1 C; 4 G; 3 T; 0 U; 0 Other;

Query Match 50.0%; Score 9; DB 1; Length 9;

Best Local Similarity 100.0%; Pred. No. 0;

Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 4 GAGGCTGTT 12  
| | | | | | | | | |  
Db 1 GAGGCTGTT 9

RESULT 5

AAA80787 ID AAA80787 standard; DNA; 8 BP.

AAA80787; AC

DT 24-NOV-2000 (first entry)

DE A. thaliana primer walking octamer SEQ ID NO: 100.

XX primer walking; octamer; primer; DNA sequencing; PCR; ss.

OS Arabidopsis thaliana.

XX US6083695-A.

PD 04-JUL-2000.

PF 21-MAY-1997; 97US-00859954.

FR 15-APR-1996; 96US-00632782.

XX (UYHO-) UNIV HOUSTON.

PA (HARD/) HARDIN S H.

PI Hardin PE, Hardin SH, Homayouni R; WPI; 2000-474852/41.

PT Sequencing an unknown DNA molecule for the polymerase chain reaction and other primer processes comprises primer walking of octamer oligonucleotides.

PS Example 8; Col 75-76; 161pp; English.

XX This invention describes a novel method for sequencing an unknown DNA molecule which comprises selecting a library primer from an octamer oligonucleotide library consisting of 48 8-bp sequences and corresponding complementary sequences, where the library primer is complementary to a known sequence adjacent to the unknown sequence or is complementary to a sequence in a known extension product. The method is useful for DNA nucleotide sequencing, in PCR, and in other processes which make use of primers. The octamers are used to identify coding sequences. Primer walking using the octamer libraries is advantageous over other sequencing methods because it does not require multiple cloning steps nor subsequent template preparations, and it is a directed and methodical approach. AAA80588-A81253 represent the octamer primers used in the primer walking method of the invention

CC walking using the octamer libraries is advantageous over other sequencing methods because it does not require multiple cloning steps nor subsequent template preparations, and it is a directed and methodical approach.

CC AAA80588-A81253 represent the octamer primers used in the primer walking method of the invention

CC method of the invention

CC Sequence 8 BP; 1 A; 1 C; 3 G; 3 T; 0 U; 0 Other;

OY 2 TTGAGGCT 9  
| | | | | | | | | |  
Db 1 TTGAGGCT 8

Search completed: December 8, 2004, 06:35:34  
Job time : 0.001 secs

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OM nucleic - nucleic search, using sw model

Run on: December 8, 2004, 06:38:09 ; Search time 0.001 Seconds  
(without alignments)  
0.972 Million cell updates/sec

Title: US-09-913-800-21

Perfect score: 18

Sequence: 1 cttgaggtctgtgagcgcac 18

Scoring table: IDENTITY NUC  
Gapop 10\_0, Gapext 0.5

Searched: 3 segs, 27 residues

Total number of hits satisfying chosen parameters: 6

Minimum DB seq length: 8  
Maximum DB seq length: 30

Post-processing: Minimum Match 0%  
Maximum Match 100%

Listing first 3 summaries

Database : rmpb21.seq:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length DB	ID	Description
1	9	50.0	9	1 US-09-989-789-2478	Sequence 2478, Ap
2	9	50.0	9	1 US-09-990-186-2478	Sequence 2478, Ap
3	9	50.0	9	1 US-09-989-994-2478	Sequence 2478, Ap

ALIGNMENTS

RESULT 1  
US-09-989-789-2478  
; Sequence 2478, Application US/09989789  
; Patent No. US20020063379A1  
; GENERAL INFORMATION:  
; APPLICANT: Liu, Qiang  
; TITLE OF INVENTION: POSITION DEPENDENT RECOGNITION OF GNN NUCLEOTIDE  
; FILE REFERENCE: 8325-0011.20 / S11-US2  
; CURRENT APPLICATION NUMBER: US/09/989,789  
; NUMBER OF SEQ ID NOS: 4085  
; SOFTWARE: PatentIn Ver. 2.0  
; SEQ ID NO 2478  
; LENGTH: 9  
; TYPE: DNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: Description of Artificial Sequence: example target  
; OTHER INFORMATION: DNA  
US-09-989-789-2478

Query Match 50.0%; Score 9; DB 1; Length 9;  
Best Local Similarity 100.0%; Pred. No. 0;  
Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 4 GAGGCTGTT 12  
Db 1 GAGGCTGTT 9

RESULT 2

US-09-990-186-2478  
; Sequence 2478, Application US/09990186  
; Publication No. US20030068675A1  
; GENERAL INFORMATION:  
; APPLICANT: Liu, Qiang  
; TITLE OF INVENTION: POSITION DEPENDENT RECOGNITION OF GNN NUCLEOTIDE  
; FILE REFERENCE: 8325-0011.21 / S11-US3  
; CURRENT APPLICATION NUMBER: US/09/990,186  
; NUMBER OF SEQ ID NOS: 4085  
; SOFTWARE: PatentIn Ver. 2.0  
; SEQ ID NO 2478  
; LENGTH: 9  
; TYPE: DNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: Description of Artificial Sequence: example target  
US-09-990-186-2478

Query Match 50.0%; Score 9; DB 1; Length 9;  
Best Local Similarity 100.0%; Pred. No. 0;  
Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 4 GAGGCTGTT 12  
Db 1 GAGGCTGTT 9

RESULT 3

US-09-989-994-2478  
; Sequence 2478, Application US/09989994  
; Publication No. US20030104526A1  
; GENERAL INFORMATION:  
; APPLICANT: Liu, Qiang  
; TITLE OF INVENTION: POSITION DEPENDENT RECOGNITION OF GNN NUCLEOTIDE  
; FILE REFERENCE: 8325-0011.20 / S11-US2  
; CURRENT APPLICATION NUMBER: US/09/989,994  
; NUMBER OF SEQ ID NOS: 4085  
; SOFTWARE: PatentIn Ver. 2.0  
; SEQ ID NO 2478  
; LENGTH: 9  
; TYPE: DNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: Description of Artificial Sequence: example target  
US-09-989-994-2478

Query Match 50.0%; Score 9; DB 1; Length 9;  
Best Local Similarity 100.0%; Pred. No. 0;  
Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 4 GAGGCTGTT 12  
Db 1 GAGGCTGTT 9

Search completed: December 8, 2004, 06:38:10  
Job time : 1 secs



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OM nucleic - nucleic search, using bw model  
Run on: December 8, 2004, 06:41:41 ; Search time 0.001 Seconds  
(without alignments)  
0.360 Million cell updates/sec

Title: US-09-913-800-32  
Perfect score: 18  
Sequence: 1 GTGAGCGACTTCATCCCT 18  
Scoring table: IDENTITY NUC  
Gapop 10.0, Gapext 0.5

Searched: 1 seqs, 10 residues  
Total number of hits satisfying chosen parameters: 2

Minimum DB seq length: 8  
Maximum DB seq length: 30  
Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 1 summaries

Database: rge32.seq:\*  
Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length DB	ID	Description
C 1	10	55.6	10 1	BD239875	ACCESSION:BD239875

ALIGNMENTS

RESULT 1  
BD239875/c 10 bp DNA Linear PAT 17-JUN-2003

LOCUS BD239875  
DEFINITION Preparation and use of superior vaccines.  
ACCESSION BD239875  
VERSION BD239875.1 GI:33049645  
KEYWORDS JP 2002534056-A/1293.  
SOURCE Homo sapiens (human)  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Cranialia; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
1 (bases 1 to 10)  
Roberts, B.L. and Shankara, S.  
Preparation and use of superior vaccines  
PATENT: JP 2002534056-A 1293 15-OCT-2002;  
GENZYME CORP  
OS Homo sapiens (human)  
PN JP 2002534056-A/1293  
PD 15-OCT-2002  
PF 18-JUN-1999 JP 2000554749  
PR 19-JUN-1998 US 60/090039, 19-JUN-1998 US 60/090040 PR  
19-JUN-1998 US 60/090041, 19-JUN-1998 US 60/089853 PR  
19-JUN-1998 US 60/089997, 19-JUN-1998 US 60/090079 PR  
19-JUN-1998 US 60/090035, 19-JUN-1998 US 60/089933 PR  
19-JUN-1998 US 60/089992, 19-JUN-1998 US 60/090072 PR  
19-JUN-1998 US 60/089878, 19-JUN-1998 US 60/089991 PR  
19-JUN-1998 US 60/090000, 19-JUN-1998 US 60/090048 PR

19-JUN-1998 US 60/089999, 19-JUN-1998 US 60/090043 PR  
19-JUN-1998 US 60/090042, 19-JUN-1998 US 60/090036 PR  
19-JUN-1998 US 60/090044, 19-JUN-1998 US 60/089844 PR  
19-JUN-1998 US 60/090080, 19-JUN-1998 US 60/089933 PR  
19-JUN-1998 US 60/089994, 19-JUN-1998 US 60/090077 PR  
19-JUN-1998 US 60/090078, 19-JUN-1998 US 60/090047 PR  
19-JUN-1998 US 60/090076, 19-JUN-1998 US 60/090045 PR  
08-DEC-1998 US 60/1111715  
PI BRUCE L ROBERTS, SRINIVAS SHANKARA  
PC C12N1/19'  
PC C12N1/21, C12N5/10, G01N33/15, G01N33/50, G01N33/53, G01N33/566, PC  
G01N37/00,  
PC C12N15/00, C12N5/00, C12N15/00  
PC Preparation and use of superior vaccines  
FH Key Location/Qualifiers  
FT source 1. .10  
FT /organism='Homo sapiens (human)'.  
FEATURES  
source 1. .10  
/organism='Homo sapiens'  
/mol\_type='genomic DNA'  
/db\_xref='taxon:9606'

Query Match 55.6%; Score 10; DB 1; Length 10;  
Best Local Similarity 100.0%; Pred. No. 0;  
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 2 TGAGCGACTT 11  
DB 10 TGAGCGACTT 1

Search completed: December 8, 2004, 06:41:41  
Job time: 0.001 secs

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OM nucleic - nucleic search, using bw model

Run on: December 8, 2004, 06:43:02 ; Search time 0.001 Seconds  
(without alignments)  
1.008 Million cell updates/sec

Title: US-09-913-800-32  
Perfect score: 18  
Sequence: 1 gtgagcgacttcctcct 18

Scoring table: IDENTITY\_NTC  
Gapop 10.0 , Gapext 0.5

Searched: 2 seqs, 28 residues

Total number of hits satisfying chosen parameters: 4  
Minimum DB seq length: 8  
Maximum DB seq length: 30

Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 2 summaries

Database : rng32.seq:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	18	100.0	18	AAA08479	Human Akt-2 phosph
2	10	55.6	10	AAZ78865	Human dendritic ce

ALIGNMENTS

RESULT 1  
AAA08479 standard; DNA; 18 BP.  
AAA08479:  
17-JUN-2000 (first entry)

Human Akt-2 phosphorothioate antisense oligonucleotide SEQ ID NO:32.  
Human; Akt-2; antisense oligonucleotide; phosphorothioate; inhibition;  
KW serine/threonine kinase; antiinflammatory; cytosolic; antiinfectious;  
KM gene therapy; infection; inflammation; tumour; ss.  
XX Homo sapiens.  
OS  
XX  
XX Key Location/Qualifiers  
FH modified\_base 1.18  
FT /\*tag= a  
FT /note= "phosphorothioate linkages"  
EN US6043090-A.  
XX  
XX 28-MAR-2000.  
XX  
XX 23-FEB-1999; 99US-00256465.

XX 23-FEB-1999; 99US-00256465.  
XX (ISIS-) ISIS PHARM INC.  
XX  
XX Monia BP, Cowsett LM;  
DR WPI; 2000-270345/23.  
XX  
XX Antisense compound for diagnosis and treatment of infection, inflammation  
PT and tumor formation is targeted towards the nucleic acid encoding a  
PT member of serine/threonine family of kinases.  
PS Claim '3; Col 38; 30pp; English.  
XX  
XX The present invention describes antisense compounds of about 8-30  
CC nucleotides in length targeted to the 5' UTR (untranslated region), 3'  
CC UTR or coding region of the nucleic acid encoding human Akt-2, which  
CC inhibits the expression of human Akt-2. Human Akt-2 is a member of the  
CC Akt/PKB family of serine/threonine kinases. The antisense compounds have  
CC antiinflammatory, cytosolic and antiinfectious activities, and can be  
CC used in gene therapy. They are useful in inhibiting the expression of  
CC human Akt-2 by contacting the cells or the tissues in vitro. They can  
CC also be used for diagnosis and treatment of infection, inflammation and  
CC tumour formation, and for prophylaxis. The present sequence represents a  
CC human Akt-2 phosphorothioate antisense oligonucleotide used in the  
CC exemplification of the present invention  
SQ Sequence 18 BP; 3 A; 5 C; 4 G; 6 T; 0 U; 0 Other;

Query Match 100.0%; Score 18; DB 1; Length 18;  
Best Local Similarity 100.0%; Pred. No. 0;  
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
Oy 1 GTGAGCGACTTCCTT 18  
Db 1 GTGAGCGACTTCCTT 18

RESULT 2  
AAZ78865/c  
ID AAZ78865 standard; DNA; 10 BP.  
AAZ78865:  
10-APR-2000 (first entry)

Human dendritic cell SAGE tag, SEQ ID NO:1293.  
XX  
XX SAGE tag; serial analysis of gene expression; antigen-presenting cell;  
KM APC; monocycle-derived dendritic cell; differential gene expression;  
KW immunostimulatory cofactor; costimulatory factor; CTL;  
KM cytotoxic T-lymphocyte; tumour antigen; immunotherapy; anticancer; ss.  
XX  
XX Homo sapiens.  
OS  
XX  
XX WO9965924-A2.  
XX  
XX 23-DEC-1999.  
XX  
XX 16-JUN-1999; 99NO-US013800.  
XX  
XX 19-JUN-1998; 98US-0089833P.  
XX 19-JUN-1998; 98US-0089844P.  
XX 19-JUN-1998; 98US-0089853P.  
XX 19-JUN-1998; 98US-0089878P.  
XX 19-JUN-1998; 98US-0089912P.  
XX 19-JUN-1998; 98US-0089929P.  
XX 19-JUN-1998; 98US-0089933P.  
XX 19-JUN-1998; 98US-0089944P.  
XX 19-JUN-1998; 98US-0089957P.  
XX 19-JUN-1998; 98US-0089959P.  
XX 19-JUN-1998; 98US-0090000P.

PR 19-JUN-1998; 98US-0090035P.  
 PR 19-JUN-1998; 98US-0090036P.  
 PR 19-JUN-1998; 98US-0090039P.  
 PR 19-JUN-1998; 98US-0090040P.  
 PR 19-JUN-1998; 98US-0090041P.  
 PR 19-JUN-1998; 98US-0090042P.  
 PR 19-JUN-1998; 98US-0090043P.  
 PR 19-JUN-1998; 98US-0090044P.  
 PR 19-JUN-1998; 98US-0090045P.  
 PR 19-JUN-1998; 98US-0090047P.  
 PR 19-JUN-1998; 98US-0090048P.  
 PR 19-JUN-1998; 98US-0090072P.  
 PR 19-JUN-1998; 98US-0090076P.  
 PR 19-JUN-1998; 98US-0090077P.  
 PR 19-JUN-1998; 98US-0090078P.  
 PR 19-JUN-1998; 98US-0090079P.  
 PR 19-JUN-1998; 98US-0090080P.  
 PR 08-DEC-1998; 98US-0111715P.

XX (GENZ ) GENZYME CORP.  
 PA (ROBE/) ROBERTS B L.  
 PA (SHAN/) SHANKARA S.

PI Roberts BL, Shankara S;

DR WPI; 2000-106077/09.

PT Isolated polynucleotides differentially expressed in antigen-presenting  
 cells, useful in gene vaccines against cancer.

XX Claim 1; Page 102; 130pp; English.

CC Sequences AA27573-279709 represent SAGE (serial analysis of gene  
 CC expression) tags used to identify mRNA transcripts encoding  
 CC immunostimulatory cofactor proteins which are preferentially or  
 CC differentially expressed in monocyte-derived dendritic cells compared  
 CC with monocytes. Some of the transcripts correspond to known genes or ESTs  
 CC (expressed sequence tags) which were previously unknown to be  
 CC preferentially or differentially expressed in dendritic cells, while  
 CC other transcripts correspond to novel genes. Antigen-presenting cell  
 CC (APC)-associated costimulatory factors play an important role in the  
 CC activation of the cytotoxic immune response, particularly against tumour  
 CC cells. Tumour antigen presentation via the MHC (major histocompatibility  
 CC complex) and subsequent recognition by T-cell receptors is alone  
 CC insufficient to activate a robust cytotoxic immune response that can lyse  
 CC the tumour cells. Immunostimulatory cofactors also being required for  
 CC efficient activation of cytotoxic T-lymphocytes (CTLs). Nucleic acid  
 CC sequences identified using the SAGE tags have several potential uses.  
 CC They may be used in vaccines to induce an immune response, particularly  
 CC against a tumour antigen; to modulate the genotype of an APC; to screen  
 CC for agents that modulate expression of differentially expressed genes in  
 CC an APC; and as hybridisation probes/amplification primers for the  
 CC diagnosis, prognosis and monitoring of diseases related to abnormal  
 CC expression of these genes. Detection of the dendritic cell differentially  
 CC expressed genes, or of their encoded proteins, can be used to identify  
 CC cells as belonging to the monocyte lineage. Cells containing these genes  
 CC can be used in active immunotherapy (or to stimulate production of a  
 CC population of antigen-specific effector cells) and vectors containing  
 CC them are used in gene therapy. Co-administration of tumour antigens and  
 CC APC-associated costimulatory factors ensures adequate antigen  
 CC presentation to endogenous APCs and upregulates the APCs for the  
 CC presentation of co-stimulatory signals, migration to T cell-rich sites,  
 CC secretion of T cell growth factors and secretion of chemokines for  
 CC recruitment of immune effector cells

CC SQ Sequence 10 BP; 3 A; 3 C; 2 G; 2 T; 0 U; 0 Other;

Query Match 55.6%; Score 10; DB 1; Length 10;  
 Best Local Similarity 100.0%; Pred. No. 0;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 2 TGAGGCACTT 11  
 |||||

DB 10 TGAGGCACTT 1  
 Search completed: December 8, 2004, 06:43:02  
 Job time : 0.001 secs



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OM nucleic - nucleic search, using sw model

Run on: December 8, 2004, 06:45:59 ; Search time 0.001 Seconds  
(without alignments)  
0.360 Million cell updates/sec

Title: US-09-913-800-32

Perfect score: 18

Sequence: 1 gtcgagcacttcaccctt 18

Scoring table: IDENTITY NUC  
Gapop 10\_0 , Gapext 0.5

Searched: 1 seqs, 10 residues

Total number of hits satisfying chosen parameters: 2

Minimum DB seq length: 8  
Maximum DB seq length: 30  
Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 1 summaries

Database : rnpb32.seq\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length DB	ID	Description
C 1	10	55.6	10	1	US-10-033-145-1293 Sequence 1293, Ap

ALIGNMENTS

RESULT 1  
 US-10-033-145-1293/C  
 ; Sequence 1293, Application US/10033145  
 ; Publication No. US20020151515A1  
 ; GENERAL INFORMATION:  
 ; APPLICANT: GENZYME CORPORATION  
 ; APPLICANT: ROBERTS, BRUCE  
 ; APPLICANT: SHANKARA, SRINIVAS  
 ; TITLE OF INVENTION: PREPARATION AND USE OF SUPERIOR VACCINES  
 ; FILE REFERENCE: GA0201C  
 ; CURRENT APPLICATION NUMBER: US/10/033,145  
 ; CURRENT FILING DATE: 2001-11-05  
 ; PRIOR APPLICATION NUMBER: PCT/US99/13800  
 ; PRIOR FILING DATE: 1999-06-18  
 ; NUMBER OF SEQ ID NOS: 2137  
 ; SOFTWARE: PatentIn version 3.0  
 ; SEQ ID NO 1293  
 ; LENGTH: 10  
 ; TYPE: DNA  
 ; ORGANISM: Homo sapiens  
 ; US-10-033-145-1293

Query Match 55.6%; Score 10; DB 1; Length 10;  
 Best Local Similarity 100.0%; Pred. No. 0;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 TGAGCGACTT 11

Db 10 TGAGCGACTT 1  
 Search completed: December 9, 2004, 06:46:00  
 Job time : 0.001 secs

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OM nucleic - nucleic search, using sw model

Run on: December 8, 2004, 06:36:40 : Search time 0.001 Seconds  
 (without alignments)  
 0.936 Million cell updates/sec

Title: US-09-913-800-21

Perfect score: 18

Sequence: 1 cttgagcctgttgccgac 18

Scoring table: IDENTITY\_NUC  
 Gapop 10.0, Gapext 0.5

Searched: 2 seqs, 26 residues

Total number of hits satisfying chosen parameters: 4

Minimum DB seq length: 8

Maximum DB seq length: 30

Post-processing: Minimum Match 0%  
 Maximum Match 100%  
 Listing first 2 summaries

Database : rni21.seq:\*  
 Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

Result No.	Score	Query Match	Length	ID	Description
1	18	100.0	18	1	US-09-256-465-21
2	8	44.4	8	1	US-08-859-954-100

ALIGNMENTS

RESULT 1  
 US-09-256-465-21  
 ; Sequence 21, Application US/09256465  
 ; Patent No. 6043090  
 ; GENERAL INFORMATION:  
 ; APPLICANT: Brett P. Monia  
 ; TITLE OF INVENTION: ANTISENSE MODULATION OF AKT-2 EXPRESSION  
 ; FILE REFERENCE: RNS-0035  
 ; CURRENT APPLICATION NUMBER: US/09/256,465  
 ; CURRENT FILING DATE: 1999-02-23  
 ; NUMBER OF SEQ ID NOS: 47  
 ; SEQ ID NO 21  
 ; LENGTH: 18  
 ; TYPE: DNA  
 ; ORGANISM: Artificial Sequence  
 ; FEATURE:  
 ; OTHER INFORMATION: Antisense Oligonucleotide  
 US-09-256-465-21

Query Match 100.0%; Score 18; DB 1; Length 18;  
 Best Local Similarity 100.0%; Pred. No. 0;  
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 CTTGAGCCTGTTGGCGAC 18  
 |||

Db 1 CTTGAGCCTGTTGGCGAC 18

RESULT 2  
 US-08-859-954-100  
 ; Sequence 100, Application US/08859954  
 ; Patent No. 6083695  
 ; GENERAL INFORMATION:  
 ; APPLICANT: Hardin, Susan H.  
 ; APPLICANT: Homayouni, Ramtin  
 ; APPLICANT: Hardin, Paul E.  
 ; TITLE OF INVENTION: Design and Optimized Primer Library for  
 ; TITLE OF INVENTION: Gene Sequencing and Method Thereof  
 ; NUMBER OF SEQUENCES: 566  
 ; CORRESPONDENCE ADDRESS:  
 ; ADDRESSEE: Fulbright & Jaworski L.L.P.  
 ; STREET: 1301 McKinney, Suite 5100  
 ; CITY: Houston  
 ; STATE: Texas  
 ; COUNTRY: U.S.A.  
 ; ZIP: 77010-1095  
 ; COMPUTER READABLE FORM:  
 ; MEDIUM TYPE: Floppy disk  
 ; COMPUTER: IBM PC compatible  
 ; OPERATING SYSTEM: PC-DOS/MS-DOS  
 ; SOFTWARE: Patent In Release #1.0, Version #1.30  
 ; CURRENT APPLICATION DATA:  
 ; APPLICATION NUMBER: US/08/859,954  
 ; FILING DATE:  
 ; CLASSIFICATION:  
 ; PRIOR APPLICATION DATA:  
 ; APPLICATION NUMBER: 08/632,782  
 ; FILING DATE:  
 ; ATTORNEY/AGENT INFORMATION:  
 ; NAME: Paul, Thomas D.  
 ; REGISTRATION NUMBER: 32,714  
 ; REFERENCE/DOCKET NUMBER: D-5900  
 ; TELECOMMUNICATION INFORMATION:  
 ; TELEPHONE: 713/651-5325  
 ; TELEFAX: 713/651-5246  
 ; INFORMATION FOR SEQ ID NO: 100:  
 ; SEQUENCE CHARACTERISTICS:  
 ; LENGTH: 8 base pairs  
 ; TYPE: nucleic acid  
 ; STRANDEDNESS: single  
 ; TOPOLOGY: linear  
 ; MOLECULE TYPE: other nucleic acid  
 ; DESCRIPTION: /desc = "Oligonucleotide"  
 ; HYPOTHEICAL: YES  
 ; ANTI-SENSE: YES  
 US-08-859-954-100

Query Match 44.4%; Score 8; DB 1; Length 8;  
 Best Local Similarity 100.0%; Pred. No. 0;  
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 2 TTGAGCCT 9  
 |||  
 Db 1 TTGAGCCT 8

Search completed: December 8, 2004, 06:36:41  
 Job time : 1 secs

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OM nucleic - nucleic search, using sw model

Run on: December 8, 2004, 06:34:24 ; Search time 0.001 Seconds  
 (without alignments)  
 0.324 Million cell updates/sec

Title: US-09-913-800-21

Perfect score: 18  
 Sequence: 1 cttgagcctgttgcgcac 18

Scoring table: IDENTITY\_NUC  
 Gapop 10.0, Gapext 0.5

Searched: 1 seqs, 9 residues

Total number of hits satisfying chosen parameters: 2

Minimum DB seq length: 8  
 Maximum DB seq length: 30

Post-processing: Minimum Match 0%  
 Maximum Match 100%  
 Listing first 1 summaries

Database : rge21.seq:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	9	50.0	9	1 AX669029	ACCESSION:AX669029

ALIGNMENTS

RESULT 1  
 AX669029  
 LOCUS AX669029 9 bp DNA Linear PAT 26-MAR-2003  
 DEFINITION Sequence 2478 from Patent WO0242459.  
 ACCESSION AX669029  
 VERSION AX669029.1 GI:29292006  
 KEYWORDS  
 SOURCE synthetic construct  
 ORGANISM synthetic construct  
 SOURCE artificial sequences.  
 REFERENCE 1  
 AUTHORS Liu,Q.  
 TITLE Position dependent recognition of gmn nucleotide triplets by zinc fingers  
 JOURNAL Patent: WO 0242459-A 2478 30-MAY-2002;  
 FEATUERS Sangamo Biosciences Inc. (US)  
 SOURCE Location/Qualifiers  
 1..9  
 /organism="synthetic construct"  
 /mol\_type="genomic DNA"  
 /db\_xref="taxon:32630"  
 /note="example target DNA"

Query Match 50.0%; Score 9; DB 1; Length 9;  
 Best Local Similarity 100.0%; Pred. No. 0;  
 Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 4 GAGCCTGT 12  
 |||||  
 Db 1 GAGCCTGT 9

Search completed: December 8, 2004, 06:34:24  
 Job time : 0.001 secs